



- prising in its genome the chimeric gene of Claim 4;  
d from the plant of Claim 5;  
ed host cell comprising the chimeric gene of Claim 4;  
med host cell of Claim 7 wherein said host cell and a microbial cell.  
de comprising all or a substantial portion of S:2, 4, 6, 8 and 10.  
or increasing methionine content of the host cells;  
rming plant cells with the chimeric gene of Claim 4;  
g fertile mature plants from the untransformed host cells under conditions suitable to obtain seeds; and  
ny seed of step (b) for those seeds of step (a) which are untransformed seeds.  
or producing plant methionine synthase;  
rming host cells with the chimeric gene of Claim 4;

(b) growing the transformed microbial cells obtained from step (a) under conditions that result in expression of a plant methionine synthase protein.

12. The method of Claim 11 wherein the host cell is a microbial cell.

13. A method for evaluating at least one compound for its ability to inhibit the activity of a plant methionine synthase, the method comprising the steps of:
- (a) transforming a host cell with a chimeric gene comprising a nucleic acid fragment encoding a plant methionine synthase, operably linked to suitable regulatory sequences;
  - (b) growing the transformed host cell under conditions that are suitable for expression of the chimeric gene wherein expression of the chimeric gene results in production of the plant methionine synthase encoded by the operably linked nucleic acid fragment in the transformed host cell;
  - (c) optionally purifying the plant methionine synthase expressed by the transformed host cell;
  - (d) treating the plant methionine synthase with a compound to be tested; and
  - (e) comparing the activity of the plant methionine synthase that has been treated with a test compound to the activity of an untreated plant methionine synthase,
- thereby selecting compounds with potential for inhibitory activity.

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